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⑰ **Method for imparting antimicrobial activity from acrylics.**

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### Description

#### Technical field

This application relates to a method for imparting non-fugitive antimicrobial activity to an article of manufacture by forming the article from acrylic polymer fibers or fabrics.

#### Background

It is known that antimicrobial chemical agents can be spun into or topically applied to synthetic fibers. However, topical treatments are readily subjected to surface abrasion and eventually removed from the fiber. Additives are fugitive and after a limited number of wash and wear cycles the chemical agent can be depleted and the antimicrobial protection lost. A key uncertainty in these approaches is the additive level. The amount of additive needed to provide desired end use properties will depend upon the lifetime of the final product, the degree of blending with other fibers, the frequency of wash-and-wear cycles and the degree of abrasion during normal usage. The combination of these factors could require excessive additive levels which become prohibitive both economically and technically.

The main problem with the spun in approach as shown for example in CH—A5—570 407 is the toxicity of the additive. In certain end uses such as children's garments, medical products and intimate apparel, fugitive additives could have potentially toxic effects. Also, many synthetic fiber processes require elevated temperatures for extended periods of time or make use of exotic solvents under heat and pressure which could lead to unwanted side reactions of the additives that produce toxic by-products (e.g., dioxins), corrosive species (e.g., chlorides) and/or eliminate the additive's antimicrobial activity. This would result in production, handling and disposal problems and would require additional technologies to safely produce the fiber.

Surface-bonded, alkoxysilane quaternary-ammonium chlorides provide antimicrobial fiber protection. The bioactive quaternary-ammonium alkoxysilane is applied onto the fiber or final product from a methanol finish solution, and bonded to the surface by hydrolysis and condensation of the alkoxysilane with hydroxyl groups of the fiber. But in the case of acrylic, polyester, nylon and other fibers which are made of polymers that do not contain significant amounts of hydroxyl groups, the alkoxysilanes cannot be bonded to the fiber surface by the hydrolysis and subsequent condensation reactions. Thus the siloxane would be merely a topical treatment to these fibers. Several problems have been encountered in this approach. The surface-applied finish has a fugitive chloride counter ion which can easily exchange with organic phosphates and sulfonates; thus when the product is laundered the antimicrobial efficacy of the surface applied finish can be greatly reduced. Another significant problem is methanol release during the alkoxysilane bonding process, which can pose toxicity and potential flammability concerns. Finally, methoxysilyl compounds are incompatible with most aqueous, emulsion-based, textile finishes, which leads to undesirable, finish-application and durability problems.

#### Summary of the invention

It has now been discovered that certain types of acrylic polymers, fibers and fabrics have non-fugitive antimicrobial properties. In the present invention it was discovered that non-fugitive antimicrobial activity is imparted to acrylic polymers, fibers or fabrics made thereof, by copolymerization of an acrylic protonated amine comonomer and/or by use of protonated amine end groups.

The protonated amine sites are part of the acrylic polymer chain and thus their durability in the final product is independent of end use properties, such as lifetime of the final product, wash-and-wear cycle frequency and degree of abrasion during normal usage. Furthermore, diffusible, toxic additives which may pose serious concern in certain end uses are not present. Since the antimicrobial agent is part of the polymer itself, toxic byproducts from spinning and fiber processing conditions do not occur.

In addition, the protonated amines are distributed throughout the polymer and thus are present in all parts of the fiber and its surface. The equilibrium distribution of protonated amines is an inherent part of the fiber. The finding of antimicrobial activity of the fiber itself is surprising in light of the fact, as set out in U.S. Patent 3,197,430, that fibers produced from acrylonitrile polymers were thought not to have the inherent ability to inhibit the growth of bacteria. The fiber of this invention is quite stable especially relative to the surface bound alkoxysilane quaternary amines which are readily deactivated by soap solutions.

According to AT—B—319 458 which relates to the preparation of polyacrylonitrile filaments of increased hot-wet modulus and whiteness by cross-linking the chains of polyacrylonitrile, one possibility of such a cross-linking structure selected from several compounds is the use of compounds containing protonated  $\text{NH}_2$  groups, but the use of protonated amines as cross-linking agent is for increasing the hot-wet modulus. One cannot draw the conclusion that protonated amine groups as part of the chain (backbone) impart antimicrobial properties to the polymer.

An essential prerequisite to antimicrobial action is effective contact between the chemical agent and the microorganism. Upon effective contact, the chemical agent interacts with some critical component of the microbial structure and/or metabolism. Specific modes of action of antimicrobial compounds are difficult to determine and are complicated by concentration, pH, temperature, solvent, electrolyte and phase separation effects.

The term antimicrobial is used to describe reduction in the number of viable pathogenic or nuisance

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causing (e.g., odor causing, biological or physical degradation causing) microorganisms. The term non-fugitive is used to describe the nature of the acrylic antimicrobial component, which, barring exposure of the acrylic polymer itself to severe conditions of hydrolysis or decomposition, cannot be eliminated from the fiber.

5 These non-fugitive antimicrobial properties of this invention are ideal for reduction of disease causing microorganisms, protection against biological degradation, and are useful for odor reduction and mildew resistance, etc.

10 Acrylonitrile polymers, fibers and fabrics with antimicrobial properties would be useful as antimicrobial air and water filters for isolation purposes, solution stabilizers and membranes for purification and concentration purposes, medical devices for disease prevention in bandages, wound dressings, wipes, tampons, sponges, gowns, drainage tubes, catheters, sutures, etc., hygiene products for odor reduction in diapers, sanitary napkins, socks, etc., industrial products for mildew resistant paints, antimicrobial packaging materials for increased fruit and vegetable shelf life and for many other uses where microorganisms are encountered.

15 Additives may also be used with this invention to impart additional antimicrobial or other hygienic, biological, or medical protection to acrylic polymers, fibers or fabrics in end uses where additional reduction in the number of viable pathogenic or nuisance causing microorganisms or elimination or deactivation of metabolic byproducts is desired.

### 20 Polymer preparation method:

One liter of demineralized water was added to a clean two liter water jacketed aluminum vessel. The water was blanketed with nitrogen gas and agitated between 50 and 150 revolutions per minute. The reaction temperature was set between 50°C and 60°C with heated water through the reaction vessel jacket. Individual monomers were weighed out in the desired amounts so the sum total of their weights was 400 25 grams. If used, the amine monomer was added to the reaction vessel first, and then the pH was adjusted with 10% sulfuric acid to  $3.0 \pm 0.5$ . The remaining monomers were carefully added to the reaction vessel. The pH was checked again and adjusted if necessary. The temperature was checked and allowed to come back into limits before addition of the initiating agents. In the meantime, one of the following polymerization initiator solutions was prepared:

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|------|--|
| (1a) | 8 grams of 77 ppm ferrous ammonium sulfate solution        |
|      | 20 grams of 11.5% hydrogen peroxide solution               |
|      | 30 grams of 7% 1-thioglycerol solution                     |
| (1b) | 8 grams of 77 ppm ferrous ammonium sulfate solution        |
| 35   | 20 grams of 11.5% hydrogen peroxide solution               |
|      | 30 grams of 7% thiourea solution                           |
| (2)  | 8 grams of 77 ppm ferrous ammonium sulfate solution        |
|      | 30 grams of 20% sodium bisulfite solution                  |
|      | 20 grams of 4.5 potassium persulfate solution              |
| 40   | (3) 5 grams of 2,2'-azobis(2-amidinopropane) hydrochloride |
|      | 45 grams of demineralized water.                           |

The initiator solution was then added to the reaction vessel, and the ensuing reaction was watched closely. The agitator speed was slowly increased as the slurry began to thicken. Room temperature 45 demineralized water was slowly added as the temperature began to rise. The reaction was run from the point of initiation for 60 minutes. If the slurry became too thick or the reaction temperature went too high, a small amount of slurry was drawn out and more room temperature demineralized water was slowly added.

At the end of the reaction time one of the following polymer quench solutions was added to the reaction vessel:

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|-----|---|
| (1) | 10 grams of 8% ethylenediaminetetraacetic acid solution   |
|     | 15 grams of 20% sodium bisulfite solution                 |
| (2) | 10 grams of 8% ethylenediaminetetraacetic acid solution   |
| 55  | (3) Cold water to reduce reaction temperature below 45°C. |

After the appropriate quench solution was added, the polymer slurry was stirred for 15 minutes. The slurry was drawn out of the reaction vessel by vacuum aspiration and filtered on a Buckner funnel. The polymer was then removed from the filter funnel, reslurried in one liter of 80°C demineralized water, stirred for 10 minutes, and filtered. The reslurry filtration step was repeated twice. The polymer was then slurried 60 in reagent grade acetone, stirred 10 minutes, and filtered on a Buckner funnel. The washed polymer was then wrapped in cheese cloth, placed in a stainless steel pan and dried 16 to 24 hours in an air draft oven at 65°C.

### Fiber preparation

65 Fibers were prepared by standard acrylic dry spinning techniques. The polymer was dissolved in

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dimethylformamide up to a 23 to 33% solids level, heated to between 90 and 110°C, filtered, then heated to 110 to 150°C and extruded through a 1400 hole spinneret at a spinning speed of 270 ypm. Nitrogen aspiration gas carried the solvent away as the individual filaments were formed. The fibers collected at the exit of the spinning cell had an as-spun denier per filament of 9.0. Residual solvent was extracted from the spun fiber in 90 to 100°C water as the fiber was drawn either 4.5× or 6.0× to yield drawn denier per filaments of 2.0 and 1.5 respectively. Standard textile fiber finish was applied using a roll applicator to reduce interfiber friction and provide lubricity during subsequent textile processing. The finish applied contained a cationic antistat rather than an anionic antistat, since the anionic antistat would react with the protonated amine. The drawn fiber was then mechanically crimped and collected at the exit end of a draw-extraction machine. The mechanically crimped yarn was then dried in a dryer between 90 and 120°C to yield finel denier per filament of 3.0 and 2.2 respectively.

### Antimicrobial activity measurement procedures:

The non-fugitive antimicrobial activity of polymers, fibers and fabrics were evaluated by the Shake  
15 Flask Test Procedure and the Parallel Streak Method.

A. Shake Flask Test—In the Shake Flask Test procedure, 70 ml of sterile pH 7.2 potassium hydrogen phosphate buffer solution was added to sterile 250 ml Erlenmeyer flasks. Each flask was then inoculated with 5 ml of a bacterial inoculum to yield 10<sup>6</sup> colony forming units/ml solutions. The flasks were shaken well and time zero counts were made by removing 1 ml aliquots from each flask and making 1:10 and 1:100  
20 dilutions in phosphate buffer. Duplicate pour plates of the 1:100 dilution in tryptone glucose extract agar were prepared, incubated at 37°C for 18 to 24 hours and counted. One gram of test polymer or fiber was aseptically added to each flask. One extra flask was prepared containing the bacterial inoculum but no test solution to ensure that the bacterial culture was viable in the test solution. The flasks were placed on a wrist  
25 action shaker and shaken for one to two hours. The one to two hour contact time counts were made by removing 1 ml aliquots from each flask and making 1:10 and 1:100 dilutions in phosphate buffer. Duplicate pour plates of both dilutions were prepared in tryptone glucose extract agar, incubated at 37°C for 18 to 24  
30 hours and the dilutions containing 30 to 300 counts were counted. The zero time counts and the one hour contact time counts were then corrected for their respective dilution factors, and 100 times the difference between the initial and final counts divided by the initial counts was the test item percent reduction. The  
shake flask procedure provided results on the relative bioactivities of various polymers, fibers and fabrics against a wide range of microorganisms. Results are given in Tables II—VI and VIII. The Shake Flask Test procedure is outlined in Malek, J. R. and Speier, J. L., *The Journal of Coated Fabrics*, Vol. 12, July, 1982, pp. 38—45.

B. Parallel Streak Method—To confirm the non-fugitivity of the antimicrobial component, the Parallel  
35 Streak Method was employed. Sterilized agar growth medium was prepared and 15 ml poured into each 100 mm diameter flat bottom petri dish. The agar was then allowed to firmly gel. A 1 ml sample of the desired test organism broth culture was transferred into 9 ml of sterile demineralized water and mixed by stirring with a 4 mm inoculating loop. One loopful of the diluted inoculum was transferred to a sterile agar  
40 plate by making five 7.5 cm long parallel streaks 1 cm apart in the center of the plate without refilling the loop, taking care not to break the agar surface. An oblong shaped test specimen was gently pressed transversely across the five inoculation streaks to insure intimate contact with the agar surface. The sample was then incubated for 18 to 24 hours at 37°C. The incubated plates were examined for interruption of growth along the streaks of inoculum underneath the sample and for a clear zone beyond the sample edge. The average width of the zone of inhibition around the test specimen was estimated with a ruler graduated  
45 in millimeters. Results are given in Table VII. This method is described in more detail in AATCC Test Method 147—1977. The polymeric acrylonitrile compositions are effective against the following microorganisms; gram-positive or gram-negative bacteria, fungi, yeasts, spore-forming bacteria, molds, staphylococcus, pseudomonas, streptococcus and bacillus.

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TABLE I  
Amine containing acrylic monomers  
and initiating agents

		M.W. g/mole
5	Monomers	
	Dimethylaminoethylmethacrylate (DMAM)	157
10	Diethylaminoethylmethacrylate (DEAM)	185
	Tertiarybutylaminoethylmethacrylate (TBAM)	185
	Dimethylaminoneopentylacrylate (DMANPA)	174
15	2-Methyl-5-vinyl pyridine (MVP)	119
	Dimethylaminopropylmethacryloamide (DMAPMA)	170
20	End groups	
	Thiourea	76
	2-Amidinopropane hydrochloride	121

## Example 1

Polymer samples were prepared according to the method described above. Acrylonitrile (AN) and methylacrylate (MA) monomers were copolymerized with various protonated amine containing monomers. These polymers were then challenged against *Klebsiella pneumoniae* in the Shake Flask Test. Results from these experiments are given in Table II. This example shows that the copolymerization of protonated amine containing monomers in acrylic polymers imparts antimicrobial activity.

TABLE II

Item	Polymerization initiator	Percent monomer by weight				Shake flask % reduction ( <i>Klebsiella</i> )	Quench system
		AN	MA	Amine	(Name)		
1	2	94	6	0	—	40.0	2
2	2	90	6	4	(DMAM)	96.9	2
3	2	90	6	4	(DEAM)	99.7	2
4	2	90	6	4	(TBAM)	99.9	2
5	2	90	6	4	(DMANPA)	95.0	2
6	2	90	6	4	(MVP)	74.3	2
7	2	90	6	4	(DMAPMA)	96.2	2

## Example 2

To determine the effects of protonated amine containing end groups on polymer bioactivity, a range of polymerization initiators were tested. Polymer samples were prepared according to the method described and challenged against *Klebsiella pneumoniae* in the Shake Flask Test. Results from these experiments, given in Table III, show that protonated amine containing end groups impart bioactivity to acrylic polymers.

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TABLE III

Item	Polymerization initiator	Percent monomer by weight		Shake flask % reduction (Klebsiella)	Quench mixture	End group*
		AN	MA			
8	1a	94	6	36.4	1	thioglycerol
9	1b	94	6	96.1	1	thiourea
10	2	94	6	40.0	2	sulfonate
11	3	94	6	99.8	3	2-amidino-propane

\* Thiourea and 2-amidino-propane end groups contain protonated amines. Thioglycerol and sulfonate end groups do not contain protonated amines.

## Example 3

To determine the minimum copolymerized protonated amine level required to produce a bioactive polymer, a series of polymers were prepared according to the method described above and then challenged against *Klebsiella pneumoniae* in the Shake Flask Test. Results from these experiments are given in Table IV.

TABLE IV

Item	Polymerization initiator	Percent monomer by weight			Shake flask % reduction (Klebsiella)	Quench mixture
		AN	MA	DEAM		
12	1a	94	6	0	36.4	1
13	1a	93.8	6	0.2	77.4	1
14	1a	93.5	6	0.5	83.1	1
15	1a	93	6	1	66.2	1
16	1a	92	6	2	98.7	1
17	1a	90	6	4	95.3	1

## Examples 4—7

Examples 4, 5, 6, 7 include Items 18 and 19. As can be seen from Table V, Item 18 does not include any protonated amine, whereas Item 19 contains a protonated amine.

## Example 4

Polymers were prepared in a continuous overflow reaction and then spun into fibers according to the method described above. The bioactivity of the polymer and the fiber were determined in the Shake Flask Test against *Klebsiella pneumoniae*. Results of these experiments are given in Table V. These results demonstrate that bioactive polymer properties are retained when transformed into fiber.

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TABLE V

Item	Polymerization initiator	Percent monomer by weight			Shake flask % reduction (Klebsiella)		Quench mixture
		AN	MA	DEAM	Polymer	Fiber	
18	2	94	6	0	2.3	3.4	2
19	1a	90.7	6	3.3	98.8	99.9	1

## Example 5

Fiber samples were prepared according to the method described above and were challenged against a wide range of microorganisms in the Shake Flask Test. Results of these experiments are given in Table VI. These results demonstrate that fibers made from protonated amine containing acrylic polymers have a high level of bioactivity against a wide range of microorganisms.

TABLE VI

Microorganism	Shake flask % reduction	
	Item 18	Item 19
A. niger (mold)	19.7	62.5
S. cerevisiae (yeast)	33.1	74.4
Cladosporium sp. (mildew)	20.1	60.7
Bacterial inoculum		
K. pneumoniae (Klebsiella)	14.5	99.9
E. coli	8.2	99.9
E. agglomerans	32.0	99.9
S. aureus	23.1	99.9
S. epidermidis	25.1	99.9
B. subtilis	49.4	85.3
P. aeruginosa	54.0	99.0

## Example 6

Fiber samples were prepared according to the method described above and challenged against Klebsiella pneumoniae in the Parallel Streak Method, to confirm the nonfugitivity of amine containing acrylic fibers. Results from these experiments are given in Table VII.

TABLE VII  
Parallel streak method

Item	Growth inhibition (Klebsiella)	
	Below sample	Zone around sample (mm)
18	None	0
19	Complete	0

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## Example 7

Fiber samples prepared by the method described above were fabricated into nonwoven products and challenged against *Klebsiella* in the Shake Flask Test. Results from these experiments are given in Table VIII. These results demonstrate that the high level non-fugitive bioactivity found in protonated amine containing acrylic fibers is maintained in nonwoven fabrics.

TABLE VIII

	Shake flask % reduction ( <i>Klebsiella</i> )	
	Item 18	Item 19
Fabric		
Hydraulically needled	38.5	99.9
Thermally bonded (25% polyester binder)	44.3	99.9

## Claims

1. A method for imparting non-fugitive antimicrobial activity, effective in reducing viable microorganisms, exhibiting a shake flask percent reduction of *Klebsiella* of 70 to 100% to an article of manufacture comprising the steps of forming said article of manufacture from a polymeric acrylonitrile composition of:
  - a. at least 85% by weight acrylonitrile,
  - b. up to about 13% by weight of a neutral ethylenically unsaturated monomer, and
  - c. from about 0.1 to 10% by weight of a protonated amine containing compound,
 where said antimicrobial activity is inherent due to protonated amine sites as part of the acrylonitrile polymer chain.
2. The method of claim 1 wherein the protonated amine containing compound is a protonated amine containing comonomer.
3. The method of claim 1 wherein the protonated amine containing compound is a protonated amine containing polymer end group.
4. The method of claim 2 wherein the protonated amine containing comonomer is an acrylic monomer selected from the group consisting of
  - dimethylaminoethylmethacrylate (DMAM),
  - diethylaminoethylmethacrylate (DEAM),
  - tertiarybutylaminoethylmethacrylate (TBAM),
  - dimethylaminoneopentyl acrylate (DMANPA),
  - 2-methyl-5-vinyl pyridine (MVP), and
  - dimethylaminopropylmethacrylamide (DMPMA).
5. The method of claim 3 wherein the protonated amine containing polymer end group is an amidino group, connected to the polymer backbone, selected from the group consisting of thiourea and 2-amidinopropane hydrochloride.
6. The method of claim 2 or 3 wherein the microorganisms are gram-positive or gram-negative bacteria.
7. The method of claim 2 or 3 wherein the microorganisms are fungi.
8. The method of claim 2 or 3 wherein the microorganisms are yeasts.
9. The method of claim 2 or 3 wherein the microorganisms are spore-forming bacteria.
10. The method of claim 2 or 3 wherein the microorganisms are molds.
11. The method of claim 2 or 3 wherein the microorganisms are staphylococcus.
12. The method of claim 2 or 3 wherein the microorganisms are pseudomonas.
13. The method of claim 2 or 3 wherein the microorganisms are streptococcus.
14. The method of claim 2 or 3 wherein the microorganisms are bacillus.
15. The method of claim 2 or 3 wherein the article is a filter that would come in contact with the microorganisms.
16. The method of claim 2 or 3 wherein the article is a medical device that would come in contact with biological fluids.
17. The method of claim 2 or 3 wherein the article is a container for packaging food.
18. The method of claim 2 or 3 wherein the shake flask percentage reduction of *Klebsiella* is 99.9—100%.
19. The method of claim 2 or 3 wherein the acrylonitrile composition is a fiber, having a finish containing cationic antistat applied thereto.



Patentansprüche

1. Verfahren, einem Fabrikationsgegenstand eine nichtflüchtige, antimikrobielle Aktivität zu verleihen, das die lebensfähigen Mikroorganismen wirksam vermindert und bei Klebsiella in der Schüttelflasche eine 70- bis 100%ige Reduktion herbeiführt, welches als Schritt die Bildung des Fabrikationsgegenstands aus einer polymeren Acrylnitrilzusammensetzung aus
  - a. wenigstens 85 Gew.-% Acrylnitril,
  - b. bis zu etwa 13 Gew.-% eines neutralen, ethylenisch ungesättigten Monomers und
  - c. etwa 0,1 bis 10 Gew.-% einer protonierten aminhaltigen Verbindung,
 umfasst, wobei die antimikrobielle Aktivität aufgrund der protonierten Aminstellen als Teil der Acrylnitrilpolymerkette inhärent vorhanden ist.
2. Verfahren nach Anspruch 1, worin die das protonierte Amin enthaltende Verbindung ein ein protoniertes Amin enthaltendes Comonomer ist.
3. Verfahren nach Anspruch 1, worin die das protonierte Amin enthaltende Verbindung eine ein protoniertes Amin enthaltende polymere Endgruppe ist.
4. Verfahren nach Anspruch 2, worin das das protonierte Amin enthaltende Comonomer ein aus der aus Dimethylaminoethylmethacrylat (DMAM), diethylaminoethylmethacrylat (DEAM), t-Butylaminoethylmethacrylat (TBAM), dimethylaminoneopentylacrylat (DMANPA), 2-Methyl-5-vinylpyridine (MVP) und dimethylaminopropylmethacryloamid (DMPMA) bestehenden Gruppe ausgewähltes Acrylmonomer ist.
5. Verfahren nach Anspruch 3, worin die das protonierte Amin enthaltende polymere Endgruppe eine mit dem Polymergerüst verbundene Amidinogruppe, ausgewählt aus der aus Thioharnstoff und 2-Amidinopropan-hydrochlorid bestehenden Gruppe, ist.
6. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen gram-positive oder gram-negative Bakterien sind.
7. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen Pilze sind.
8. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen Hefen sind.
9. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen sporenbildende Bakterien sind.
10. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen Schimmelpilze sind.
11. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen Staphylokokken sind.
12. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen Pseudomonen sind.
13. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen Streptokokken sind.
14. Verfahren nach Anspruch 2 oder 3, worin die Mikroorganismen Bazillen sind.
15. Verfahren nach Anspruch 2 oder 3, worin der Gegenstand ein Filter ist, der mit Mikroorganismen in Berührung kommt.
16. Verfahren nach Anspruch 2 oder 3, worin der Gegenstand eine medizinische Vorrichtung ist, der mit biologischen Flüssigkeiten in Berührung kommt.
17. Verfahren nach Anspruch 2 oder 3, worin der Gegenstand ein Behälter zum Verpacken von Lebensmitteln ist.
18. Verfahren nach Anspruch 2 oder 3, worin die prozentuale Reduktion von Klebsiella in der Schüttelflasche 99,9 bis 100% beträgt.
19. Verfahren nach Anspruch 2 oder 3, worin die Acrylnitrilzusammensetzung eine Faser mit einer Appretur ist, die ein darauf angewandtes kationisches Antistatistikmittel enthält.

Revendications

1. Un procédé pour conférer à un article manufacturé une activité antimicrobienne persistante, efficace pour réduire des micro-organismes viables, présentant un pourcentage de réduction de Klebsiella de 70 à 100% dans l'essai en fiole secouée, comprenant les étapes de formation dudit article manufacturé à partir d'une composition polymère à base d'acrylonitrile constituée de:
  - a. au moins 85% en poids d'acrylonitrile,
  - b. au plus environ 13% en poids d'un monomère neutre éthyléniquement insaturé, et
  - c. environ 0,1 à 10% en poids d'un composé à fonction amine protonée,
 où ladite activité antimicrobienne est intrinsèque du fait que des sites aminos protonés font partie de la chaîne de polymère d'acrylonitrile.
2. Le procédé de la revendication 1, dans lequel le composé à fonction amine protonée est un comonomère à fonction amine protonée.
3. Le procédé de la revendication 1, dans lequel le composé à fonction amine protonée est un groupe terminal de polymère à fonction amine protonée.
4. Le procédé de la revendication 2, dans lequel le comonomère à fonction amine protonée est un monomère acrylique choisi dans le groupe formé par
  - le méthacrylate de diméthylaminoéthyle (MDMA),
  - le méthacrylate de diéthylaminoéthyle (MDEA),
  - le méthacrylate de tertibutylaminoéthyle (MTBA),
  - l'acrylate de diméthylaminonéopentyle (ADMANP),

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la 2-méthyl-5-vinylpyridine (MVP), et

le diméthylaminopropylméthacrylamide (DMPMA).

5 5. Le procédé de la revendication 3, dans lequel le groupe terminal de polymère à fonction amine protonée est un groupe amidino, relié au squelette du polymère, choisi dans le groupe formé par la thiourée et le chlorhydrate de 2-amidinopropane.

6. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des bactéries Gram-positives ou Gram-négatives.

7. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des champignons.

8. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des levures.

10 9. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des bactéries sporogènes.

10. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des moisissures.

11. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des staphylocoques.

12. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des Pseudomonas.

15 13. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des streptocoques.

14. Le procédé de la revendication 2 ou 3, dans lequel les micro-organismes sont des bacilles.

15. Le procédé de la revendication 2 ou 3, dans lequel l'article est un filtre qui doit entrer en contact avec les micro-organismes.

16. Le procédé de la revendication 2 ou 3, dans lequel l'article est un dispositif médical qui doit entrer en contact avec des fluides biologiques.

17. Le procédé de la revendication 2 ou 3, dans lequel l'article est un récipient pour emballer un produit alimentaire.

18. Le procédé de la revendication 2 ou 3, dans lequel le pourcentage de réduction de Klebsiella dans l'essai en fiole secouée est de 99,9 à 100%.

25 19. Le procédé de la revendication 2 ou 3, dans lequel la composition à base d'acrylonitrile est une fibre à laquelle est appliqué on apprêt contenant un agent antistatique cationique.

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